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(cushingoid syndrome), except for emergency therapy.

- (v) Clinical and experimental data have demonstrated that corticosteroids administered orally or by injection to animals may induce the first stage of parturition when administered during the last trimester of pregnancy and may precipitate premature parturition followed by dystocia, fetal death, retained placenta, and metritis.
- (vi) Federal law restricts this drug to use by or on the order of a licensed veterinarian. A withdrawal period has not been established for this product in preruminating calves. Do not use in calves to be processed for veal.
- (b)(1) Specifications. Each tablet contains 0.25 milligram of dexamethasone.¹
- (2) Sponsors. See Nos. 000061 and 061623 in §510.600(c) of this chapter.
- (3) Conditions of use—(i) Amount. Dogs: Administer orally at 0.25 to 1.25 milligrams per day for up to 7 days. Cats: 0.125 to 0.5 milligram per day for up to 7 days.
- (ii) *Indications for use*. In treatment of dogs and cats as an anti-inflammatory agent.¹
- (iii) Limitations. (a) Clinical and experimental data have demonstrated that corticosteriods administered orally or by injection to animals may induce the first stage of parturition when administered during the last trimester of pregnancy; and they may precipitate premature parturition followed by dystocia, fetal death, retained placenta, and metritis.
- (b) Do not use in viral infections. Anti-inflammatory action of corticosteroids may mask signs of infections. Do not use in animals with tuberculosis, chronic nephritis, cushingoid syndrome, or peptic ulcers, except for emergency therapy.¹
- (c) Federal law restricts this drug to use by or on the order of a licensed veterinarian.¹

[40 FR 26273, June 23, 1975, as amended at 44 FR 7130, Feb. 6, 1979; 50 FR 49372, Dec. 2, 1985; 52 FR 7832, Mar. 13, 1987; 55 FR 8461, Mar. 8, 1990; 66 FR 14073, Mar. 9, 2001; 68 FR 4914, Jan. 31, 2003; 70 FR 16934, Apr. 4, 2005]

§ 520.540c Dexamethasone chewable tablets.

- (a) Specifications. Each half-scored tablet contains 0.25 milligram of dexamethasone.
- (b) Sponsor. See No. 000069 in $\S510.600$ (c) of this chapter.
- (c) Conditions of use—(1) Amount. 0.25 to 1.25 milligrams per day.¹
- (2) *Indications for use*. Supportive therapy in nonspecific dermatosis and inflammatory conditions in dogs.¹
- (3) Limitations. (i) Administer by free-choice feeding or crumble over food. Administer 0.25 to 1.25 milligrams daily in single or two divided doses until response is noted or 7 days have elapsed. When response is attained, dosage should be gradually reduced by 0.125 milligram per day until maintenance level is achieved.
- (ii) Clinical and experimental data have demonstrated that corticosteriods administered orally or parenterally to animals may induce the first stage of parturition when administered during the last trimester of pregnancy; and they may precipitate premature parturition followed by dystocia, fetal death, retained placenta, and metritis.
- (iii) Do not use in viral infections. Anti-inflammatory action of corticosteriods may mask signs of infection. Do not use in animals with tuberculosis, chronic nephritis, cushingoid syndrome, or peptic ulcers, except for emergency therapy.¹
- (iv) Federal law restricts this drug to use by or on the order of a licensed veterinarian.

[44 FR 7130, Feb. 6, 1979, as amended at 56 FR 50653, Oct. 8, 1991; 60 FR 55659, Nov. 2, 1995]

§ 520.550 Glucose/glycine/electrolyte.

- (a) Specifications. The product is distributed in packets each of which contains the following ingredients: Sodium chloride 8.82 grams, potassium phosphate 4.20 grams, citric acid anhydrous 0.5 gram, potassium citrate 0.12 gram, aminoacetic acid (glycine) 6.36 grams, and glucose 44.0 grams.
- (b) Sponsor. See No. 000069 in $\S510.600$ (c) of this chapter.
- (c) Conditions of use. (1) Glucose/gly-cine/electrolyte is indicated for use in the control of dehydration associated with diarrhea (scours) in calves. It is used as an early treatment at the first

¹These conditions are NAS/NRC reviewed and deemed effective. Applications for these uses need not include effectiveness data as specified by §514.111 of this chapter.